Application Serial Number 10/579219 Response to Office Action dated 01/11/2008

612-455-3801

Amendments to the Claims:

This listing of claims will replace all prior versions and listings of claims in the application.

Listing of Claims:

- 1. 5. (Cancelled)
- 6. (Currently amended) Derivatives of compounds Garcinol and Isogarcinol of

$$R_1$$
 O O R_3

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respectively, wherein R1, R2 and R3, substituents of Garcinol, and R4 and R5, substituents of Isogarcinol, are selected from a group comprising consisting of O-Methoxy, O-Ethoxy, O-Isopropoxy, O-Allyloxy, O-Butoxy, O-t-Butoxy, O-Pentoxy, O-Hexyloxy, O-CH2-COOH, O-CO-CH2-CLCI, O-SO2-CH3, and 0-O-CH2-CHOH-CH3.

A process for preparation of derivatives of compound 7. (Currently amended) gGarcinol orand Isogarcinol of formula I and II, respectively, said process comprising steps of reacting gGarcinol or Isogarcinol with halo compounds to obtain the derivatives. with the selected substituents of R1, R2, R3, R4 and R5, at temperature ranging between

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- 30 40°C under alkaline conditions in presence of organic solvents, and purifying followed by purification to obtain the derivatives.
- 8. (Currently Amended) AThe process for proparation as claimed in claim 7, wherein the reacting process is carrying the reaction carried in presence of at least one of alkaline hydroxides or alkaline carbonates.
- 9. (Currently Amended) AThe process for preparation as claimed in claim 7, wherein the compounds Garcinol and Isogarcinol are in equimolar concentration.
- 10. (Currently Amended) A<u>The</u> process for preparation as claimed in claim 7, wherein the organic solvent is selected from a group comprising consisting of acetone, chloroform, MDC and EDC.
- 11. (Currently Amended) A<u>The</u> process for preparation as claimed in claim 7, wherein the purifying process of the derivatives are purified conducted by column chromatograpy.
- 12. (Currently Amended) A method of treating a-diseases condition selected from a group comprising cancer, asthma, cardiac hypertrophy, Acquired Immunodeficiency Syndrome (AIDS), Human Immunodeficiency-Virus (HTV) caused by histone acctyltransferase (HAT) in a subject in need thereof, wherein said method comprises a step of administering a pharmaceutically effective amount of the derivatives of compounds Garcinol or Isogarcinol of

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respectively to the subject, wherein R1, R2 and R3, substituents of Garcinol, and R4 and R5, substituents of Isogarcinol, are selected from a group comprising consisting of O-Methoxy, O-Ethoxy, O-Isopropoxy, O-Allyloxy, O-Butoxy, O-t-Butoxy, O-Pentoxy, O-Hexyloxy, O-CH2-COOH, O-CO-CH2-CLCl, O-SO2-CH3, and O-O-CH2-CHOH-CH3 to the subject.

- 13. (Currently Amended) AThe method as claimed in claim 12, wherein the derivatives are histone acetyl transferase (HAT) inhibitors.
- 14. (New) The process as claimed in claim 7, wherein said halo compounds are selected from a group consisting of halogens and HOCO-CH₂-Cl.

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15. (New) The method as claimed in claim 12, wherein the diseases caused by HAT are at least one selected from a group consisting of cancer, asthma, cardiac hypertrophy and acquired immunodeficiency syndrome.